

**AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of claims

- 1-13. (Cancelled)
14. (Currently amended) An *in vitro* method for producing mature dendritic Langerhans cells, said method comprising:
- a. culturing cells selected from the group consisting of peripheral blood monocytes and bone marrow cells in a medium containing mammalian platelets; and
  - b. incubating the culture at about 30°C to about 40°C for a period sufficient to enable formation of mature dendritic Langerhans cells, wherein the medium omits an exogenous cytokine, wherein the platelets and the peripheral blood monocytes or bone marrow cells are derived from the same species or from phylogenetically close species.
15. (Previously presented) The method as claimed in claim 14, wherein the exogenous cytokine is granulocyte macrophage colony stimulating factor or interleukin-4.
16. (Previously presented) The method as claimed in claim 14 wherein the medium comprises RPMI-1640.
17. (Previously presented) The method as claimed in claim 14 wherein the cells are cultured for a period of about 2 to about 8 days.
18. (Previously presented) The method as claimed in claim 14 wherein the medium further comprises fetal calf serum.

19. (Previously presented) The method as claimed in claim 18, wherein the medium contains at least about 2% fetal calf serum.
20. (Previously presented) The method as claimed in claim 19, wherein the fetal calf serum is about 10%.
21. (Currently amended) The method as claimed in claim 14 wherein human platelets are added to the medium to develop mature dendritic Langerhans cells.
22. (Cancelled)
23. (Withdrawn - Currently amended) A method for producing mature dendritic Langerhans cells *in vitro* comprising:
  - a. preparing peripheral blood monocytes and/or bone marrow cells; and
  - b. culturing the peripheral blood monocytes or the bone marrow cells with platelets of the same species in a culture medium lacking an exogenous cytokine such that mature dendritic Langerhans cells are produced.
24. (Withdrawn) The method of claim 23 further comprising analyzing the morphology of human dendritic Langerhans cells produced.
25. (Withdrawn - Currently amended) The method of claim 23 further comprising analyzing the mature dendritic Langerhans cells produced by flow cytometry.
26. (Withdrawn - Currently amended) The method of claim 23 wherein the peripheral blood monocytes, the platelets, and the mature dendritic Langerhans cells produced are human.
27. (Cancelled)
28. (Previously presented) An *in vitro* method for producing mature dendritic Langerhans cells, said method comprising:

- a. culturing cells selected from the group consisting of peripheral blood monocytes and bone marrow cells in a medium containing mammalian platelets; and
- b. incubating the culture at about 30°C to about 40°C for a period sufficient to enable formation of mature dendritic Langerhans cells,  
wherein the platelets and the peripheral blood monocytes or bone marrow cells are derived from the same species or from phylogenetically close species.

29. (Withdrawn - Currently amended) A method for producing mature dendritic

Langerhans cells *in vitro* comprising:

- a. preparing peripheral blood monocytes and/or bone marrow cells; and
- b. culturing the peripheral blood monocytes or the bone marrow cells with platelets of the same species in a culture medium such that mature dendritic Langerhans cells are produced.

30. (Withdrawn - Currently amended) A method for producing mature dendritic

Langerhans cells *in vitro* comprising:

- a. preparing peripheral blood monocytes and/or bone marrow cells; and
- b. culturing the peripheral blood monocytes or the bone marrow cells with platelets in a culture medium such that mature dendritic Langerhans cells are produced,

wherein the peripheral blood monocytes and/or bone marrow cells and the platelets may be independently selected from the group of rat cells and mouse cells.

31. (Withdrawn) The method of claim 30, wherein the culture medium lacks an exogenous cytokine.
32. (Currently amended) A method for producing mature human dendritic Langerhans cells *in vitro* comprising:
- a. preparing human peripheral blood monocytes and/or human bone marrow cells; and
  - b. culturing the human peripheral blood monocytes or the human bone marrow cells with human platelets ~~of the same species~~ in a culture medium such that mature human dendritic Langerhans cells are produced, wherein more than about 50% of the mature human dendritic Langerhans cells have dendritic processes and display reactivity to anti-HLA-DR, anti-CD40, and anti-CD86 monoclonal antibodies and less than about 20% of the mature human dendritic Langerhans cells display reactivity to anti-CD1a, anti-CD80, and anti-CD83 monoclonal antibodies.
33. (Previously presented) The method of claim 32, wherein the culture medium lacks an exogenous cytokine.
34. (Withdrawn) The method of claim 23 wherein the peripheral blood monocytes are human peripheral blood monocytes, the platelets are human platelets, and the dendritic Langerhans cells produced are human dendritic Langerhans cells.
35. (New) The method of claim 14 or 28 wherein the platelets and the peripheral blood monocytes or bone marrow cells are derived from the same species.
36. (New) The method of claim 35 wherein the same species is human.
37. (New) The method of claim 14 or 28 wherein the platelets and the peripheral blood monocytes or bone marrow cells are derived from phylogenetically close species.

38. (New) The method of claim 37 wherein the platelets are from rat and peripheral  
blood monocytes or bone marrow cells are from mouse.